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(FILE 'HOME' ENTERED AT 07:28:06 ON 04 AUG 2008)  
FILE 'CA' ENTERED AT 07:28:40 ON 04 AUG 2008  
L1 5564 S (COLD OR CRYOGENIC) (3A) (TRAP? OR FINGER OR CAPTUR?) OR CRYOTRAP?  
L2 33 S L1 AND(STAGE# OR STEP? OR SEQUEN?) (4A) (HEAT? OR DESOR? OR  
RELEASES?)  
L3 63 S L1 AND TRAP?(4A) (ISOLAT? OR EVACUAT?)  
L4 21 S L1 AND TRAP?(4A)VACUUM AND VALVE  
L5 117 S L2-4  
L6 105 S L5 AND PY<2003

=> d bib,ab 16 1-105

L6 ANSWER 5 OF 105 CA COPYRIGHT 2008 ACS on STN  
AN 137:244108 CA  
TI On-line coupled superheated water extraction (SWE) and superheated water chromatography (SWC)  
AU Tajuddin, Ruziyati; Smith, Roger M.  
CS Department of Chemistry, Loughborough University, Loughborough, Leicestershire, LE11 3TU, UK  
SO Analyst (Cambridge, United Kingdom) (2002), 127(7), 883-885  
AB Superheated water extn. has been linked directly to a superheated water chromatog. sepn. so that the process of sample extn. and sepn. can be achieved without the need for org. solvents at any stage. A model matrix spiked with pharmaceuticals and antioxidants was extd. and the extd. components were collected on a cold polystyrene-divinylbenzene trap. The analytes were then sequentially released by raising the temp. in stages. Each fraction was passed online to a polystyrene divinylbenzene anal. column and was eluted with superheated water using a thermal gradient.

L6 ANSWER 28 OF 105 CA COPYRIGHT 2008 ACS on STN  
AN 126:347069 CA  
OREF 126:67413a,67416a  
TI Determination of butyltin compounds in sediments by means of hydride generation/cold trapping gas chromatography coupled to inductively coupled plasma mass spectrometric detection  
AU Garcia, E. Segovia; Alonso, J. I. Garcia; Sanz-Medel, A.  
CS Department of Physical and Analytical Chemistry, Faculty of Chemistry, Oviedo, 33006, Spain  
SO Journal of Mass Spectrometry (1997), 32(5), 542-549  
AB A method for the detn. of butyltin compds. in sediments is based on the generation of volatile mono-, di- and tributyltin (MBT, DBT, TBT) hydrides from a 4% (vol./vol.) acetic acid medium using NaBH4. The hydrides formed are then trapped on a Chromosorb W HP SP2100 packed glass column immersed in liq. N. Sequential desorption of the hydrides is achieved by Nichrome wire heating of the column. The MBT, DBT and TBT hydrides are detected by mass spectrometry using an inductively coupled plasma source. Detection limits were 7, 4 and 4 pg (as Sn) for MBT, DBT and TBT, resp. The method was applied to the detn. of organotin compds. (DBT and TBT) in the certified ref. material CRM 462 with satisfactory results.

